

| REF       | 90231024 |
|-----------|----------|
| Pack Size | 24 Cards |

# Forward Grouping card (8 Column)

| + 4°C -+ 25°C<br>+ 4°C       | Manufacturer           | IVD  In vitro Diagnostic Medical Device | Batch Number / Lot Number | Expiry date                | FG8C  |
|------------------------------|------------------------|---|---------------------------|----------------------------|---|
| Consult Instructions for use | Date of<br>Manufacture | REF Catalogue Number                    | This side up              | Keep Away<br>from Sunlight | SBIOCAT <sup>™</sup> Forward Grouping card (8 Column) |

#### SUMMARY

According to ABO blood group system human red blood cell antigen can be divided into four groups A, B, AB and O depending on the presence or absence of corresponding antigens on the red blood cells. Also human red blood cells are classified as Rho (D) positive or Rho (D) negative depending upon the presence or absence of Rho (D) antigen. It is important to examine previous transfusion and testing records in pre transfusion compatibility testing. Similarly current ABO and Rho (D) testing results should be compared with previous testing records. Concurrence between both the results gives confirmation of testing results.

## **REAGENTS**

SBIOCAT<sup>™</sup> Forward Grouping Card contains eight microtubes prefilled with a gel in a suitable buffer containing Monoclonal Anti-A (Clone 11H5), Anti-B (Clone 6F9) and Anti-D (IgM) (VI-) (Clone P3x61 + TH-28) from microtube 1 to 3 and 5 to 7. Microtubes 4 and 8 contain neutral gel for test control. SBIOCAT<sup>™</sup> Forward Grouping Card is used for the ABO grouping and Rho (D) typing.

#### STORAGE AND STABILITY

Store SBIOCAT<sup>TM</sup> gel cards in an upright position at 4-25°C. Do not freeze. Avoid exposure of SBIOCAT<sup>TM</sup> gel cards to direct sunlight or any heat source. The shelf life of SBIOCAT<sup>TM</sup> gel cards is as per the expiry date mentioned on the label. Do not use beyond expiry date. Once the aluminium foil is removed from the microtube, it should be used immediately.

# ADDITIONAL REAGENTS AND MATERIALS REQUIRED

SBIOCAT<sup>™</sup> Diluent -2 LISS for preparation of red cell suspension. (Refer package insert before use). Gel card centrifuge (85g), Work station, Micropipette capable of delivering 5-50µl of specimen and Bottle top dispenser.

#### **PRINCIPLE**

As the SBIOCAT  $^{\text{TM}}$  gel card containing red blood cells is centrifuged under specific conditions, the red blood cells possessing the corresponding antigen will agglutinate in presence of the specific antibody and will be trapped in the gel column. The red blood cells, which do not react are not trapped in the gel column and get settled at the bottom of the microtube. The reactions are then read and graded according to their reactivity pattern.

# SAMPLE COLLECTION

No special preparation of the patient is required prior to sample collection by approved techniques. For optimal results, freshly collected sample should be used. Anticoagulants like EDTA, CPD-A and Citrate can be used.

#### SAMPLE PREPARATION

Prepare a 5 % red blood cell suspension in SBIOCAT<sup>™</sup> Diluent- 2 LISS as follows:

- Bring the SBIOCAT<sup>™</sup> Diluent- 2 LISS to room temperature before testing.
- Dispense 0.5 ml of SBIOCAT™ Diluent- 2 LISS into a clean test tube.
- Add 50µl of whole blood or 25µl of packed red cells and mix gently.
- Red blood cell suspension so obtained should be used for forward grouping.

## .TEST PROCEDURE

- Label the "SBIOCAT Forward Grouping Card" with patient's name or identification number. Remove the aluminium foil of required number of microtubes carefully by pulling it backwards.
- Pipette 10µI of 5% patient's red blood cell suspension to the microtubes labeled as A-B-D-Ctrl, taking care to ensure that micropipette tip does not touches the microtube.
- 3. Centrifuge the cards for 10 minutes in the gel card centrifuge.
- 4. Retrieve the card from centrifuge, read and record the results. **Note:** For applications on SBIOCAT  $^{\text{TM}}$  HEXA, 50 $\mu$ I of 0.8-1.0% red cell suspension can be used instead of 10 $\mu$ I of 5% red cell suspension.

# INTERPRETATION OF RESULTS

The control microtube (Ctrl) must be negative to validate the forward grouping results. If not, then repeat the test after washing the patient's red blood cells with warm saline.

Positive reaction: Agglutinated red blood cells forming a clear line on the surface of gel column or agglutinates dispersed in the gel

**Negative reaction:** Non agglutinated red blood cells settle at the bottom of the microtube forming a compact button.

The reaction strength may be recorded as follows:

| Strength of reaction | Comments   |
|----------------------|--|
| 4+                   | Agglutinated red blood cells form a line on the surface of the gel microtube.    |
| 3+                   | Most agglutinated red blood cells remain in the upper half of the gel microtube. |
| 2+                   | Agglutinated red blood cells are observed throughout the length of the           |

|                              | microtube. A small button of red blood cells may also be visible at the bottom of the gel microtube.  |
|------------------------------|---|
| 1+                           | Most agglutinated red blood cells remain in the lower half of the gel microtube. A button of cells may also be visible at the bottom of the gel microtube.        |
| ±                            | Most agglutinated red blood cells are in the lower third part of the gel microtube.   |
| Negative                     | All the red blood cells pass through and form a compact button at the bottom of the gel microtube.  |
| Mixed field<br>agglutination | Agglutinated red blood cells form a line on the surface of the gel and non-agglutinated red blood cells form a compact button at the bottom of the gel microtube. |
| Н                            | Hemolysis of red blood cells  |

Note: Visual reading of reactions in a card may differ from the reactions read by any automated software through image processing. However this may not change the final result interpretation

#### Expected reactivity pattern for ABO grouping:

| Anti-A   | Anti-B   | Blood Group |
|----------|----------|-------------|
| ± to 4+  | Negative | А           |
| Negative | ± to 4+  | В           |
| ± to 4+  | ± to 4+  | AB          |
| Negative | Negative | 0           |

Note: Human red blood cells that show weak reaction with Anti-A and/or Anti-B probably indicate subgroups of A and/or B and further testing is recommended.

# Expected reactivity pattern for Rho (D) typing:

| Anti-D   | Rho(D) Type      |
|----------|------------------|
| ± to 4+  | Rho (D) Positive |
| Negative | Rho (D) Negative |

Note: Weak D/ Partial D type human red blood cells may give a weaker or negative reaction. Such cells should be retested for weak D confirmation with SBIOCAT<sup>™</sup> Coombs Anti-IgG card.

## NOTE

- In vitro diagnostic reagent for laboratory and professional use only. Not for medicinal use. The SBIOCAT  $^{\!\!\top\!\!}$  gel cards contains sodium azide
- The SDROCAT geradiate. Avoid contact with skin and mucosa. On disposal flush with large quantity of water.

  All SBIOCAT™ gel cards should be centrifuged for one complete cycle (10 minutes) in gel card centrifuge before
- Visually inspect the SBIOCAT<sup>™</sup> gel cards before use. SBIOCAT<sup>™</sup> gel cards having bubble(s) ent
- gel cards having bubble(s) entrapped within the gel can be centrifuged for two complete cycles in gel card centrifuge to remove the bubble, if bubbles are not removed the card should not be used.
- 6. SBIOCAT™ gel cards that exhibit any signs of drying (i.e. absence or reduced level of reagent buffer above the gel column), decreased volume of gel, cracked gel

- ${\tt SBIOCAT^{TM}}$  gel cards with damaged aluminium foil seal should not be used. 7.
- Freezing of SBIOCAT™ gel cards or evaporation of gel or reagent buffer due to exposure to heat may lead to erroneous results.
- Fibrin or particulate matter if present in the sample may lead to erroneous results.
- Fibrin if present in the sample may trap red blood cells on the surface of the gel column presenting a pink line. To avoid, samples should be well centrifuged at 1500g for 10. minutes before taking serum or plasma and RBCs should be washed if not collected properly in an anticoagulant.
- Use of red blood cells concentration/ volume and reagents other than those described may lead to erroneous results. Follow the instructions carefully.
- Aged or stored red blood cells may exhibit weaker reactivity than freshly collected cells.
- Old cell panels may give an unclear background with SBIOCAT<sup>™</sup> gel cards.
- Do not use hemolysed samples.

  Extreme turbidity or discoloration may indicate microbial contamination or denaturation of protein due to thermal damage. Such SBIOCAT™ gel cards should be discarded
- Contamination of reagents during usage may cause
- false positive or negative results.
  Red cell aggregation in the red cell suspension may interfere the passage.
- Aluminium foil seal of SBIOCAT™ gel cards should be removed gently and carefully by pulling the foil seal backwards to avoid contamination of reagents from one microtube to another.
- Do not use lipemic. icteric and hyperproteic samples
- To avoid contamination always use fresh tips before
- dispensing into each microtube.

  SBIOCAT™ ContaVoid (Cat. No. 903300100) can be used to avoid contamination of reagents in microtubes while usage.

#### REMARKS

- Known positive and negative control should be tested as per Good Laboratory Practices.
- SBIOCAT™ Red Cell Preserving Solution (Cat. No.90262020) can be used as red blood cell preservative solution for preservation of known cells.
- The Anti-D does not detect the DVI variant.

#### **BIBLIOGRAPHY**

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11 Yishun Street 51, 304-23, The Criterion, Singapore 767971.